

LABORATORIES



PHARMACEUTICALS

D-U-N-S® NUMBER: 87-219-9445

AN ISO 9001:2008 COMPANY



Certified Impurity Reference Standards Manufacturer & Suppliers
Link for list of Impurity & COA : www.veepholabs.com



VENKAT SHINDE

Managing Director & CEO

Venkat have over 18 years of experience in the Pharmaceutical Industry as successful professional. He has worked in various Pharmaceutical top companies as head A-R&D and he was responsible for analytical support to filing ANDA and DMF, Lab compliances, analytical equipment, formulation process, impurity preparation, impurity characterization, and Unknown Impurity structure elucidation.

Within short time he brings both Veeprho Laboratories and Veeprho Pharmaceuticals S.R.O. (EU) company at respectable level in the world market and now it is admire company in the impurity standards business in all the front. His vision is to bring both companies in top 5 in the world next few years.

ABOUT VEEPRO

VEEPRHO LABORATORIES PVT LTD is an emerging, research-based global pharmaceutical company with a diverse combination of skills, resources, and capabilities that provide a specific scientific support to industry in rapidly changing healthcare environment. Company has made good examples by providing most critical impurity standards to our clients, they have huge confidence on Veeprho and Veeprho is top company in impurity standard business because of quality material, quality service and quality documentation.



COMPANY'S FOCUS ON

- Impurity Isolation from API or Drug Products by Preparative HPLC.
- Synthesis of Impurities/Metabolites.
- Compliance Analytical Laboratories for Stability Studies (Exhibit & Commercial Batches) & Analytical Method Validation

SERVICES

Our team keeps on reviewing latest pharmacopeia monograph and also keeps the impurity standards ready with certification to save users time.

IMPURITY ISOLATION

Isolation and purification of impurity from mg to g scale. We have executed most critical isolation and purification project in recent past, e.g isolation of unknown impurity from drug product and APIs and Structure elucidation.

IMPURITY SYNTHESIS

We have synthesized thousands of impurity as on date. VEEPRHO offer custom synthesis of compounds on request.



CERTIFICATION OF IMPURITY STANDARDS

All impurity standards will be supplied with certificates of analysis

- Identification by $^1\text{H-NMR}$ and Mass spectroscopy.
- Purity test by either HPLC or GC.
- Other test like ^{13}C NMR, TGA, CHN, IR will be provided on request.

All impurity COA uploaded on our web site www.veeprholabs.com

CUSTOM SYNTHESIS mg to g

VEEPRO offers world class service in chemical synthesis of a wide variety of organic compounds from milligram to kilogram scale at competitive price.

- Multi step Organic Synthesis.
- Asymmetric Synthesis using Chiral Auxiliary, Catalysis, & Resolution Techniques.
- Metal-Mediated Reactions.
- Hydrogenation.
- Toxic Chemistry.
- Peptide Synthesis.



UNKNOWN IMPURITY STRUCTURE ELUCIDATION

After isolation and purification of unknown impurity from APIs and Drug products and it is be analysed by LC - MS, HPLC, H NMR , ¹³C NMR, FT-IR and elemental analysis (CHNS) to confirm its structure.

INDIA
VEEPRHO LABORATORIES PVT LTD



CZECH REPUBLIC, (EU)
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